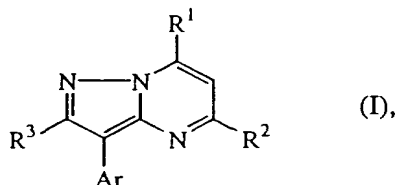


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ABSTRACT

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PYRAZOLOPYRIMIDINES  
AS CRF RECEPTOR ANTAGONISTS

This invention concerns compounds of formula



including the stereoisomers and the pharmaceutically acceptable acid addition salt forms thereof, wherein  $R^1$  is  $NR^4R^5$  or  $OR^5$ ;  $R^2$  is  $C_{1-6}$ alkyl,  $C_{1-6}$ alkyloxy or  $C_{1-6}$ alkylthio;  $R^3$  is hydrogen,  $C_{1-6}$ alkyl,  $C_{1-6}$ alkylsulfonyl,  $C_{1-6}$ alkylsulfoxy or  $C_{1-6}$ alkylthio;  $R^4$  is hydrogen,  $C_{1-6}$ alkyl, mono- or di( $C_{3-6}$ cycloalkyl)methyl,  $C_{3-6}$ cycloalkyl,  $C_{3-6}$ alkenyl, hydroxy $C_{1-6}$ alkyl,  $C_{1-6}$ alkylcarbonyloxy $C_{1-6}$ alkyl or  $C_{1-6}$ alkyloxy $C_{1-6}$ alkyl;  $R^5$  is  $C_{1-8}$ alkyl, mono- or di( $C_{3-6}$ cycloalkyl)methyl,  $Ar^1CH_2$ ,  $C_{1-6}$ alkyloxy $C_{1-6}$ alkyl, hydroxy $C_{1-6}$ alkyl,  $C_{3-6}$ alkenyl, thienylmethyl, furanylmethyl,  $C_{1-6}$ alkylthio $C_{1-6}$ alkyl, morpholinyl, mono- or di( $C_{1-6}$ alkyl)amino $C_{1-6}$ alkyl, di( $C_{1-6}$ alkyl)amino,  $C_{1-6}$ alkylcarbonyl $C_{1-6}$ alkyl,  $C_{1-6}$ alkyl substituted with imidazolyl; or a radical of formula  $-Alk-O-CO-Ar^1$ ; or  $R^4$  and  $R^5$  taken together with the nitrogen atom to which they are attached may form an optionally substituted pyrrolidinyl, piperidinyl, homopiperidinyl or morpholinyl group; having CRF receptor antagonistic properties; pharmaceutical compositions containing such compounds as active ingredients; methods of treating disorders related to hypersecretion of CRF such as depression, anxiety, substance abuse, by administering an effective amount of a compound of formula (I).